

May 31, 2013

Document Processing Desk 6(a)(2)
Office of Pesticide Programs – 7504P
U.S. Environmental Protection Agency
Ariel Ross Building
1200 Pennsylvania Avenue, NW
Washington, D.C. 20460-0001

RE: Section 6(a)(2) March Incident Filing

Dear 6(a)(2) Administrator:

On behalf of Reckitt Benckiser, Scientific & Regulatory Consultants, Inc. (SRC) is submitting the enclosed documents containing alleged adverse effect incidents for products listed below. SafetyCall is a primary gathering source for incidents, though internal reports for calls/correspondence received directly at Reckitt Benckiser are also included. SRC is acting on Reckitt Benckiser's behalf by assisting them in registration actions and their reporting requirements in accordance with FIFRA section 6(a)(2).

The EPA Registration Numbers with adverse effect incidents for this filing are:

- 777-71
- 3282-81
- 777-81
- 777-72

These incidents are being reported in compliance with 40 CFR § 159.184 and have been assigned the H-C severity classification. If additional information is needed, please feel free to contact us by e-mail (bmacdonald@srcconsultants.com) or by phone at 260-244-6270.

Sincerely,

Bob MacDonald Consultant (SRC)

Agent for Reckitt Benckiser

Bah Marchald

ann M. Kline

Consultant (SRC)
Agent for Reckitt Benckiser

P.O. Box 1014 Columbia City, IN 46725

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Phone: 260-244-6270 Fax: 260-244-6273

Personal privacy information

Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area

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Row I Administrative	Reporter Name		Submission date.	Contact person	n (if different than reporter)	Internal ID 1143539	
Data	Address Dallas, TX 75217 USA			Address			
	Phone #		Phone #				
	Incident Status: New			Date registran became aware incident. 04/03/2013		Was incident part of larger study? No	
Row 2 Pesticide(s) Involved	EPA Registration # (Product 1) 3282-81		EPA Registration # (Product 2)		EPA Registration	# (Product 3)	
	A.I. (s)		A.I. (s)		A.I. (s)	A.I. (s)	
	Product 1 name d-CON Ready Mixed Baitbits		Product 2 Name		Product 3 Name		
	Exposed to concentrate prior to dilution? No		Exposed to concentrate prior to dilution?		Exposed to concer dilution?	Exposed to concentrate prior to dilution?	
	Formulation solid		Formulation		Formulation		
Row 3 Incident Circumstances	Evidence label directions were not followed? Yes Intentional misuse? Yes Applicator certified? UNK	Incident site: (examples incluyard, school, industrial, nursery/greenhouse, surface w commercial turf, building/offiwoods, agricultural (specify oway (rail, utility, highway)). Own Residence		vater, ce, forest/	include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/		
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) See Incident Description Notes			DATE	REVIEWED FOR 6(a)2 5.24.13 INITIALS:	AK	

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Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 2 of 3

Brief description of incident circumstances.

Bryan, Kyle Apr 3 2013 3:21PM

H: The caller states his brother had a seizure earlier today and the paramedics came and treated his brother. He states his brother uses illegal drugs but the reason he is calling is because his brother ingested most of the contents of one box of the product 10 hrs ago. He denies any other sxs.

Yeager, Greg Apr 4 2013 10:32AM

CB complete. Brother has had no issues related to product ingestion, and is being treated for other problems. Passed on information on product to MD to call with any questions.

If any new or unexpected symptoms develop, please contact us 24/7 and refer to your reference number so that we can advise on further treatment.



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Demographic information: Age: 34 Year(s) Sex: Male Occupation (if relevant) NA	Exposure route: Ingestion/oral	Was adverse effect result of suicide/homicide or attempted suicide/homicide? Yes	Was protective clothing worn (specify)? None Reported
If female, pregnant? NA	Was exposure occupational? Not indicated If yes, days lost due to illness: NA	Time between exposure and onset of symptoms: Unable to determine	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). ER/Hospital-Unknown disposition Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown	List signs/symptoms/adverse eff Neurological-Seizure (single)	ects	If lab tests were performed, list test names and results (If available, submit reports) None Reported
Human severity category: HC			
necessary)		rmation surrounding the incident.	
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			Internal ID # 0 1143539 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0